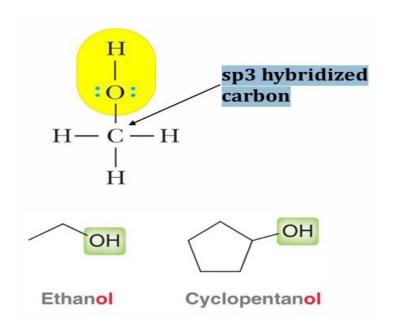
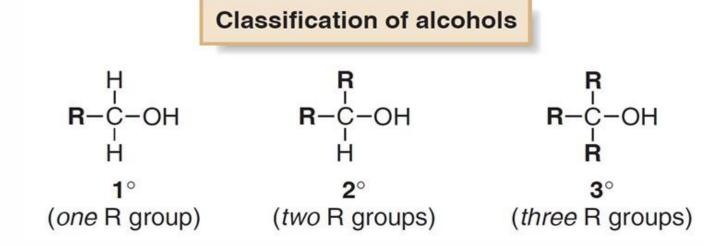
Alcohols are compounds that possess a hydroxyl group (OH) connected to an sp<sup>3</sup> hybridized carbon atom, and are characterized by names ending in "OL":



Alcohols are important because they can be converted sp3 hybridized carbon into many other types of compounds, including alkenes, haloalkanes, aldehydes, ketones, carboxylic acids, and esters. Not only can alcohols be converted to these compounds, but these compounds can also be converted to alcohols. Thus, alcohols play a central role in the interconversion of organic functional groups

### **CLASSIFICATION OF ALCOHOLS**

Alcohols are classified as primary (1°), secondary (2°), or tertiary (3°) based on the number of carbon atoms bonded to the carbon with the OH group.



**Primary alcohols** (methanol, ethanol, propanol, butanol, etc.) exhibit antibacterial properties.

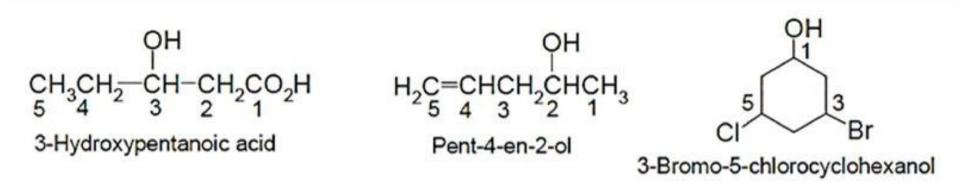
### Nomenclature of alcohols

Generally, the name of an alcohol ends with-ol. An alcohol can be named as an alkyl alcohol, usually for small alkyl groups e.g. methyl alcohol and ethyl alcohol. The longest carbon chain bearing the \_\_OH group is used as the parent; the last-e from this alkane is replaced by an-ol to obtain the root name. The longest chain is numbered starting from the end nearest to the \_\_\_ OH group, and the position of the \_\_OH group is numbered. Cyclic alcohols have the prefix cyclo-, and the \_\_OH group is deemed to be on C-1.

HO\_CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>

1-Propylcyclopentanol

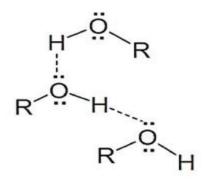
Alcohols with double or triple bonds are named using the-ol suffix on the alkene or alkyne name. Numbering gives the hydroxyl group the lowest possible number. When numbers are also given for the multiple bond position, the position of the hydroxyl can be written immediately before the-ol prefix. If the hydroxyl group is only a minor part of the structure, it may be named as a hydroxysubstituent



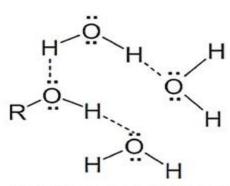
Diols are compounds with two hydroxyl groups. They are named as for alcohols except that the suffix-diol is used and two numbers are required to locate the hydroxyls. 1,2-diols are called glycols. The common names for glycols usually arise from the name of the alkene from which they are prepared.

## Physical properties of alcohols

Alcohols can be considered as organic analogues of water. Both the C\_O and O\_H bonds are polarized due to the electronegativity of the oxygen atom. The highly polar nature of the O\_H bond results in the formation of hydrogen bonds with other alcohol molecules or other hydrogen bonding systems, e.g. water and amines. Thus, alcohols have considerably higher boiling point due to the hydrogen bonding between molecules (intermolecular hydrogen bonding). They are more polar than hydrocarbons, and are better solvents for polar molecules.



Hydrogen bonds between alcohol molecules



Hydrogen bonds to water in aqueous solution

# Preparation of alcohols

## 1. Hydration of Alkenes

$$egin{aligned} H_2O+H^+ &
ightarrow H_3O^+ \ &
ightarrow C=C < +H_2O \leftrightarrow 
ightarrow CH-C-OH < \ &
ightarrow CH_3CH=CH_2+H_2O \leftrightarrow CH_3CH-OHCH_3 \end{aligned}$$

## 2. Hydroboration of Alkenes

$$CH_3CH = CH_2 + (H - BH_2)_2 \rightarrow CH_3CH_2CH_2BH_2 \longrightarrow (CH_3CH_2CH_2)_2BH$$
 
$$(CH_3CH_2CH_2)_2BH \rightarrow (CH_3CH_2CH_2)_3B \longrightarrow 3CH_3 - CH_2 - CH_2 - OH + B(OH)_3$$

## 3. Hydroformylation of Alkenes

$$R^{1}$$
 $R^{2}$ 
 $R^{3}$ 
 $CO/H_{2}$ 
 $R^{1}$ 
 $R^{3}$ 
 $R^{2}$ 
 $R^{3}$ 
 $R^{2}$ 
 $R^{3}$ 
 $R^{4}$ 
 $R^{3}$ 
 $R^{4}$ 
 $R^{5}$ 
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 $R^{5}$ 
 $R^{5}$ 
 $R^{5}$ 

## 4. Grignard Synthesis

$$\begin{array}{c} \text{OMgX} \\ \text{H}_3\text{C}-\text{C}-\text{H} + \text{R}-\text{MgX} \longrightarrow \text{H}_3\text{C}-\text{C}-\text{H} \xrightarrow{\text{H}_2\text{O}} \text{H}_3\text{C}-\text{C}-\text{H} \\ \\ \text{Ethanal} \end{array}$$
 Secondary alcohol

# REACTIONS of ALCOHOLS

#### Acid-Catalyzed Dehydration

#### E2 Dehydration with POCI3

#### Reaction with Hydrogen Halides

#### Oxidation of Alcohols

Secondary alcohols [0] ketones

Conversion to Chlorides with SDC1z; PC13; or PC15

Conversion to Bromides with PBr3; PBr5; or PPh2Br3

Oxidative Cleavage of Vicinal Diols with HIO4